

DEC 6 2005

81/3

K053115

**Special 510(k) Summary of Safety and Effectiveness:
Line Extension to the Xia[®] Spinal System**

Submission Information

Name and Address of the Sponsor of the 510(k) Submission: Stryker Spine
2 Pearl Court
Allendale, NJ 07401

Contact Person: Simona Voic
Regulatory Affairs Project Manager
(201) 760 - 8145

Date of Summary Preparation: October 31, 2005

Device Identification

Proprietary Name: Xia[®] Spinal System

Common Name: Spinal Fixation Appliances

Classification Name and Reference: Spinal Interlaminar Fixation Orthosis,
21 CFR §888.3050
Spinal Intervertebral Body Fixation Orthosis
21 CFR §888.3060
Pedicle Screw Spinal System
21 CFR §888.3070

Predicate Device Information:

Stryker Spine Xia[®] Stainless Steel System
(K012870 & K031090)
Stryker Spine Xia[®] Spinal System (K013823)
DePuy Inc., Moss Miami Spinal System (K962628)

Predicate Device Identification

The Stryker Spine Xia[®] Spinal Fixation System consists of Monoaxial and Polyaxial Screws, Washer, Hooks, Blocker, Rods, Staples, and Connectors. The components are manufactured from either Titanium material (Ti alloy and CP Titanium) or Stainless Steel.

Description of Device Modification

This submission is intended to address a line extension to Xia® Spinal System. The line extension includes the addition of Titanium alloy and Stainless Steel 4.0 mm diameter spinal screws.

Intended Use:

The Xia® Spinal System is intended for use in the noncervical spine. When used as a pedicle screw fixation system, the Xia® Spinal System is intended for patients: (a) having severe spondylolisthesis (Grades 3 and 4) at the fifth lumbar - first sacral (L5-S1) vertebral joint; (b) who are receiving fusions using autogenous bone graft only; (c) who are having the device fixed or attached to the lumbar and sacral spine; and (d) who are having the device removed after the development of a solid fusion mass.

When used as a pedicle screw fixation system, the Xia® Spinal System is also intended to provide immobilization and stabilization of spinal segments in skeletally mature patients as an adjunct to fusion in the treatment of the following acute and chronic instabilities or deformities of the thoracic, lumbar, and sacral spine: degenerative spondylolisthesis with objective evidence of neurologic impairment, fracture, dislocation, scoliosis, kyphosis, spinal tumor, and failed previous fusion (pseudarthrosis).

When used as an anterior screw fixation system or a posterior hook and sacral/iliac screw fixation system, the Xia® Spinal System is indicated for patients with degenerative disc disease which is defined as back pain of discogenic origin with degeneration of the disc confirmed by history and radiographic studies, spondylolisthesis, fracture, spinal stenosis, spinal deformities such as scoliosis, kyphosis, lordosis, tumor, pseudoarthrosis or revision of failed fusion attempts.

The 6mm diameter rods from the DIAPASON™ Spinal System and OPUS™ Spinal System are intended to be used with the other components of the Xia® Titanium Spinal System. The Titanium Multi-Axial Cross Connectors are intended to be used with the other components of the Xia® Titanium Spinal System.

Statement of Technological Comparison:

The subject components share the same intended use, material, and basic design concepts as that of the predicate device: Stryker Spine Xia[®] Spinal System (K013823, K012870, and K031090). Mechanical testing also demonstrated comparable mechanical properties to the predicate device: Stryker Spine Xia[®] Spinal System (K013823) and DePuy Inc. Moss Miami Spinal System (K962628).



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

DEC 6 2005

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

Ms. Simona Voic
Regulatory Affairs Project Manager
Stryker Spine
2 Pearl Court
Allendale, New Jersey 07401

Re: K053115
Trade/Device Name: Xia® Spinal System
Regulation Number: 21 CFR 888.3070
Regulation Name: Pedicle screw spinal system
Regulatory Class: II
Product Code: MNI, MNH, KWQ, KWP
Dated: November 4, 2005
Received: November 7, 2005

Dear Ms. Voic:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

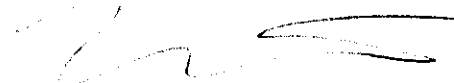
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

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This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at (240) 276-0120. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>

Sincerely yours,



SM Mark N. Melkerson
Acting Director
Division of General, Restorative
and Neurological Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

K053115

Indications for Use

510(k) Number (if known): K053115

Device Name: Xia® Spinal System

Indications For Use:

The Xia® Spinal System is intended for use in the noncervical spine. When used as a pedicle screw fixation system, the Xia® Spinal System is intended for patients: (a) having severe spondylolisthesis (Grades 3 and 4) at the fifth lumbar - first sacral (L5-S1) vertebral joint; (b) who are receiving fusions using autogenous bone graft only; (c) who are having the device fixed or attached to the lumbar and sacral spine; and (d) who are having the device removed after the development of a solid fusion mass.

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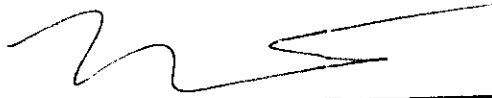
Prescription Use X
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use _____
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)

**Division of General, Restorative,
and Neurological Devices**

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